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Store at -20C  
#3247

## Pim-1 (C93F2) Rabbit mAb

For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W	<b>Reactivity:</b> H M	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 34	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P11309	<b>Entrez-Gene Id:</b> 5292
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### Product Usage Information

#### Application

Western Blotting

#### Dilution

1:1000

### Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

### Specificity/Sensitivity

Pim-1 (C93F2) Rabbit mAb detects endogenous levels of total Pim-1 protein. No cross reactivity was detected with other Pim family members.

### Source / Purification

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val160 of Pim-1.

### Background

Pim proteins (Pim-1, Pim-2 and Pim-3) are oncogene-encoded serine/threonine kinases (1). Pim-1, a serine/threonine kinase highly expressed in hematopoietic cells, plays a critical role in the transduction of mitogenic signals and is rapidly induced by a variety of growth factors and cytokines (1-4). Pim-1 cooperates with c-Myc in lymphoid cell transformation and protects cells from growth factor withdrawal and genotoxic stress-induced apoptosis (5,6). Pim-1 also enhances the transcriptional activity of c-Myb through direct phosphorylation within the c-Myb DNA binding domain as well as phosphorylation of the transcriptional coactivator p100 (7,8). Hypermutations of the Pim-1 gene are found in B-cell diffuse large cell lymphomas (9). Phosphorylation of Pim-1 at Tyr218 by Etk occurs following IL-6 stimulation and correlates with an increase in Pim-1 activity (10). Various Pim substrates have been identified; Bad is phosphorylated by both Pim-1 and Pim-2 at Ser112 and this phosphorylation reverses Bad-induced cell apoptosis (11,12). The corresponding pim-1 gene encodes a pair of proteins through use of different translation initiation sites. Both larger 44 kDa (Pim-1L) and smaller 33 kDa (Pim-1S) proteins are active kinases, but differ in stability (13).

### Background References

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5. Möröy, T. et al. (1993) *Proc Natl Acad Sci USA* 90, 10734-8.
6. Lilly, M. and Kraft, A. (1997) *Cancer Res* 57, 5348-55.
7. Levenson, J.D. et al. (1998) *Mol Cell* 2, 417-25.
8. Winn, L.M. et al. (2003) *Cell Cycle* 2, 258-62.
9. Pasqualucci, L. et al. (2001) *Nature* 412, 341-6.
10. Kim, O. et al. (2004) *Oncogene* 23, 1838-44.
11. Aho, T.L. et al. (2004) *FEBS Lett* 571, 43-9.
12. Yan, B. et al. (2003) *J Biol Chem* 278, 45358-67.
13. Saris, C.J. et al. (1991) *EMBO J* 10, 655-64.

### Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

### Western Blot Buffer

**IMPORTANT:** For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

### Applications Key

**W:** Western Blotting

### Cross-Reactivity Key

**H:** Human **M:** Mouse

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